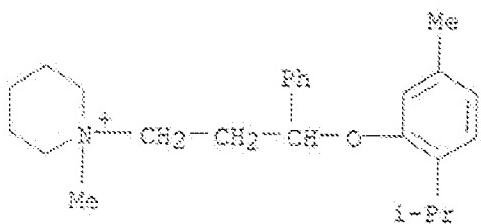


AN 1977:189458 CARPLUS  
 DM 86:189458  
 TI Aromatic amino ether quaternary ammonium salts  
 IN Ogawa, Shuntaro; Morita, Kan; Yoshida, Akiyoshi  
 PA Rohto Pharmaceutical Co., Ltd., Japan  
 SO Japan., 9 pp.  
 CODEN: JAXXAD  
 DT Patent  
 LA Japanese  
 FAN.CNT 1

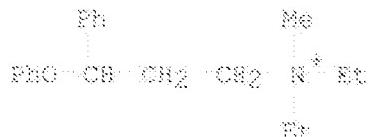
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 51044934	S4	19761201	JP 1969-99283	19691209
AB	RC <sub>6</sub> H <sub>4</sub> ZCHPhCH <sub>2</sub> CH <sub>2</sub> N <sup>+</sup> R <sub>1</sub> R <sub>2</sub> R <sub>3</sub> X <sup>-</sup> (I; R = H, halo, Me; R <sub>1</sub> , R <sub>2</sub> = H, alkyl, alkylene contg. optional O atom; R <sub>3</sub> = alkyl; Z = halo; Z = O, S) were prep'd. by quaternization of RC <sub>6</sub> H <sub>4</sub> ZCHPhCH <sub>2</sub> CH <sub>2</sub> NR <sub>1</sub> R <sub>2</sub> (II) with R <sub>3</sub> X. I were useful as antispasmodics, anticholinergics, antiinflammants, and analgesics. Thus, excess MeI was added to II [R = H, R <sub>1</sub> R <sub>2</sub> = (CH <sub>2</sub> ) <sub>5</sub> , Z = O], obtained from 2.4 g of its HCl salt after treatment with aq. NaOH and Et <sub>2</sub> O extn., in MeOH at room temp. to give 2.6 g I [R = H, R <sub>1</sub> R <sub>2</sub> = (CH <sub>2</sub> ) <sub>5</sub> , R <sub>3</sub> = Me, X = iodo, Z = O], which had anticholinergic activity with ED <sub>50</sub> of 1.6 .times. 10 <sup>-6</sup> g/mL in guinea pigs. Similarly prep'd. were 17 addnl. I and their biol. activity given.				
IT	42063-78-5P	RL: SPN (Synthetic preparation); PREP: (Preparation) (prepn. of)			
RN	42063-78-5	CARPLUS			
CN	Piperidinium, 1-methyl-1-[3-{5-methyl-2-(1-methylethyl)phenoxy}-3-phenylpropyl]-, iodide (9CI)	(CA INDEX NAME)			



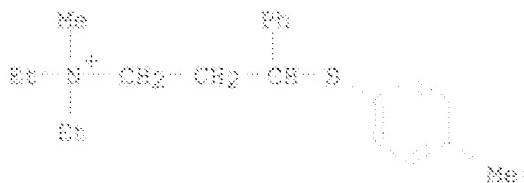
● I<sup>-</sup>

AN 1977:189458 CAPLUS  
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 TI Aromatic amino ether quaternary ammonium salts  
 IN Ogawa, Shuntaro; Morita, Ken; Yoshida, Akiyoshi  
 PA Roho Pharmaceutical Co., Ltd., Japan  
 SO Japan., 9 pp.  
 CODEN: JAXXAD  
 CT Patent  
 LA Japanese  
 IC C07C093-12  
 CC 23-4 (Noncondensed Aromatic Compounds)  
 Section-cross-reference(s): 27  
 FAN,CONT I

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 51044934	R4	19761201	JP 1969-99283	19691209
AB	RC <sub>6</sub> H <sub>4</sub> ZCHPhCH <sub>2</sub> CH <sub>2</sub> N+RIR <sub>2</sub> R <sub>3</sub> X- (I; R = H, halo, Me; R <sub>1</sub> , R <sub>2</sub> = H, alkyl, RIR <sub>2</sub> = alkylene contg. optional O atom; R <sub>3</sub> = alkyl; Z = halo; Z = O, S) were prepd. by quaternization of RC <sub>6</sub> H <sub>4</sub> ZCHPhCH <sub>2</sub> CH <sub>2</sub> NR <sub>1</sub> R <sub>2</sub> (III) with RX <sub>2</sub> . I were useful as antispasmodics, anticholinergics, antiinflammatories, and analgesics. Thus, excess MeI was added to III (R = H, RIR <sub>2</sub> = (CH <sub>2</sub> ) <sub>5</sub> , Z = O), obtained from 2.4 g of its HCl salt after treatment with sq. NaOH and Et <sub>2</sub> O extrn., in MeOH at room temp, to give 2.6 g I (R = H, RIR <sub>2</sub> = (CH <sub>2</sub> ) <sub>5</sub> , R <sub>3</sub> = Me, X = Iodo, Z = O), which had anticholinergic activity with ED <sub>50</sub> of 1.6 + 10 <sup>-8</sup> g/mL in quines' pigs. Similarly prepd. were 17 addnl. I and their biol. activity given.				
ST	quaternary arom ether antispasmodic; anticholinergic quaternary aryloxypropylammonium halide; antiinflammatory quaternary aryloxypropylammonium halide; analgesic quaternary aryloxypropylammonium halide; aryloxypropylammonium halide antispasmodic anticholinergic; aryloxypropylpiperidinium halide antispasmodic anticholinergic				
IT	Analgesics Inflammation inhibitors Muscle relaxants and Spasmolytics Parasympatholytics (aryloxypropylammonium halides)				
TT	42064-71-1P 42064-72-2P 42064-73-3P 42064-74-4P 42064-76-6P 42064-79-9P 42064-85-7P 42796-63-4P 42796-67-8P 42988-33-0P 51074-51-2P 51543-53-4P 62663-36-9P RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. and anticholinergic activity of)				
TT	42063-78-3P 42796-62-3P 42796-71-4P 43213-26-1P 51543-52-3P 62663-50-7P RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. of)				
IT	42064-87-9P 42064-89-1P RL: SPN (Synthetic preparation); PREP (Preparation) (prepn., antispasmodic and analgesic activity of)				
IT	42796-29-2 62663-37-0 62663-38-1 62663-39-2 62663-40-5 62663-41-6 62663-42-7 62663-43-8 62663-44-9 62663-45-0 62663-46-1 62663-47-2 62663-48-3 62663-49-4 RL: RCT (Reactant) (quaternization of)				
TT	42796-63-4P 62663-36-9P RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. and anticholinergic activity of)				
RN	42796-63-4 CAPLUS				
CN	Benzene propanaminium, N,N-diethyl-N-methyl-γ-phenoxy-, iodide (SCI) (CA INDEX NAME)				



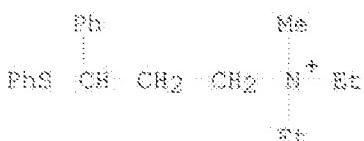
RN 62663-36-9 CAPIUS  
 CN Benzenepropanaminium, N,N-diethyl-N-methyl-gamma-((4-methylphenyl)thio)-, iodide (9CI) (CA INDEX NAME)



IT 42796-62-3P S1543-52-3P  
 BL: SPN (Synthetic preparation); PREP (Preparation)  
 (prepa. of)  
 RN 42796-62-3 CAPIUS  
 CN Benzenepropanaminium, N,N,N-trimethyl-gamma-phenoxy-, iodide (9CI) (CA INDEX NAME)

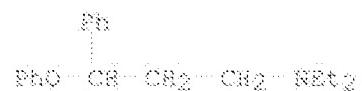


RN S1543-52-3 CAPIUS  
 CN Benzenepropanaminium, N,N-diethyl-N-methyl-gamma-(phenylthio)-, iodide (9CI) (CA INDEX NAME)



IT 62663-42-7 62663-43-8

RL: RCT (Reactant)  
(quaternization of)  
RN: 62663-42-7 CASLUS  
CN: Benzenepropanamine, N,N-diethyl- $\gamma$ -phenoxy- (9CI) (CA INDEX NAME)



RN: 62663-43-8 CASLUS  
CN: Benzenepropanamine, N,N-diethyl- $\gamma$ -((4-methylphenyl)thio)- (9CI) (CA INDEX NAME)

